

10/797,626

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NEWS X25 X.25 communication option no longer available

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=> fil reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

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TSCA INFORMATION NOW CURRENT THROUGH June 30, 2006

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REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

=> S STQNASLLSLTVC/SQSP

L1 1 STQNASLLSLTVC/SQSP

=> FIL HCAP

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	28.89	29.10

FILE 'HCAPLUS' ENTERED AT 15:22:34 ON 16 DEC 2006
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T.S. Heard Ph.D.

10/797,626

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NEWS	1		Web Page URLs for STN Seminar Schedule - N. America
NEWS	2		"Ask CAS" for self-help around the clock
NEWS	3	AUG 09	INSPEC enhanced with 1898-1968 archive
NEWS	4	AUG 28	ADISCTI Reloaded and Enhanced
NEWS	5	AUG 30	CA(SM)/CAPLUS(SM) Austrian patent law changes
NEWS	6	SEP 11	CA/CAPLUS enhanced with more pre-1907 records
NEWS	7	SEP 21	CA/CAPLUS fields enhanced with simultaneous left and right truncation
NEWS	8	SEP 25	CA(SM)/CAPLUS(SM) display of CA Lexicon enhanced
NEWS	9	SEP 25	CAS REGISTRY(SM) no longer includes Concord 3D coordinates
NEWS	10	SEP 25	CAS REGISTRY(SM) updated with amino acid codes for pyrrolysine
NEWS	11	SEP 28	CEABA-VTB classification code fields reloaded with new classification scheme
NEWS	12	OCT 19	LOGOFF HOLD duration extended to 120 minutes
NEWS	13	OCT 19	E-mail format enhanced
NEWS	14	OCT 23	Option to turn off MARPAT highlighting enhancements available
NEWS	15	OCT 23	CAS Registry Number crossover limit increased to 300,000 in multiple databases
NEWS	16	OCT 23	The Derwent World Patents Index suite of databases on STN has been enhanced and reloaded
NEWS	17	OCT 30	CHEMLIST enhanced with new search and display field
NEWS	18	NOV 03	JAPIO enhanced with IPC 8 features and functionality
NEWS	19	NOV 10	CA/CAPLUS F-Term thesaurus enhanced
NEWS	20	NOV 10	STN Express with Discover! free maintenance release Version 8.01c now available
NEWS	21	NOV 13	CA/CAPLUS pre-1967 chemical substance index entries enhanced with preparation role
NEWS	22	NOV 20	CAS Registry Number crossover limit increased to 300,000 in additional databases
NEWS	23	NOV 20	CA/CAPLUS to MARPAT accession number crossover limit increased to 50,000
NEWS	24	DEC 01	CAS REGISTRY updated with new ambiguity codes
NEWS	25	DEC 11	CAS REGISTRY chemical nomenclature enhanced
NEWS	26	DEC 14	WPIDS/WPINDEX/WPIX manual codes updated
NEWS	27	DEC 14	GBFULL and FRFULL enhanced with IPC 8 features and functionality
NEWS EXPRESS			NOVEMBER 10 CURRENT WINDOWS VERSION IS V8.01c, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006.
NEWS HOURS			STN Operating Hours Plus Help Desk Availability
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T.S. Heard Ph.D.

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FILE COVERS 1907 - 16 Dec 2006 VOL 145 ISS 26
FILE LAST UPDATED: 15 Dec 2006 (20061215/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> S L1

L2 2 L1

=> D L2 1-2 IBIB ABS HITSTR

L2 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2006:977981 HCAPLUS

DOCUMENT NUMBER: 145:334147

TITLE: Methods of inhibiting angiogenesis and tumor development

INVENTOR(S): Brooks, Peter, C.; Akalu, Abebe; Cretu, Alexandra; Policarpio, Desiree

PATENT ASSIGNEE(S): New York University, USA

SOURCE: PCT Int. Appl., 153pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006098987	A2	20060921	WO 2006-US8266	20060309
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
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US 2006216237	A1	20060928	US 2006-372367	20060309
US 2006240002	A1	20061026	US 2006-371626	20060309
PRIORITY APPLN. INFO.:			US 2005-660713P	P 20050311
			US 2005-660889P	P 20050311

US 2005-660903P P 20050311
 US 2005-711049P P 20050824
 US 2005-711177P P 20050825

AB The authors disclose methods for identifying genes and proteins modulated by antagonism of extracellular matrix (ECM) ligands that specifically interact with $\alpha\text{v}\beta 3$ integrin. The authors also disclose using the identified genes and proteins for inhibiting angiogenesis, tumor metastasis, and other tumor developmental processes, including cell migration, cell adhesion, cell proliferation, and tumor growth and for treating angiogenesis-dependent conditions. In one example, a monoclonal antibody antagonist of $\alpha\text{v}\beta 3$ is shown to modulate the expression of IGFBP-4, TSP-1, Id-1, p27KIP, and p21CIP.

IT 771528-84-8

RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

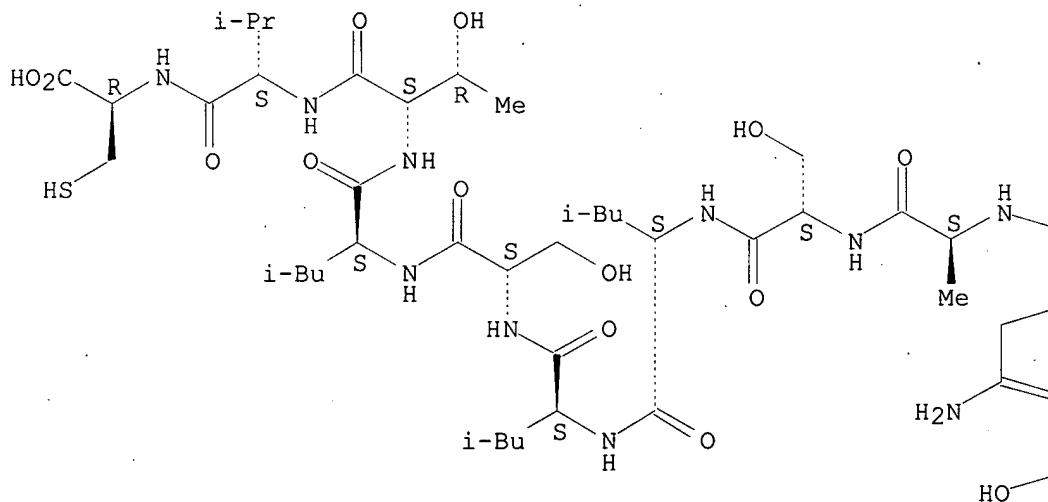
(antagonists of extracellular matrix ligand/ $\alpha\text{v}\beta 3$ integrin interaction for inhibition of tumor angiogenesis and metastasis)

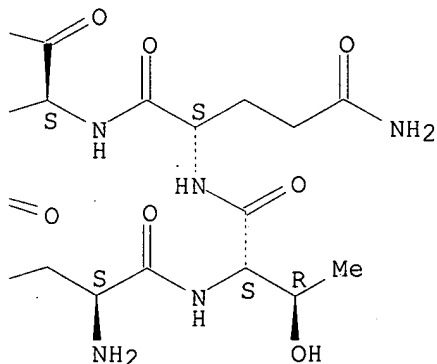
RN 771528-84-8 HCAPLUS

CN L-Cysteine, L-seryl-L-threonyl-L-glutaminyl-L-asparaginyl-L-alanyl-L-seryl-L-leucyl-L-leucyl-L-seryl-L-leucyl-L-threonyl-L-valyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A





L2 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2004:857618 HCAPLUS
DOCUMENT NUMBER: 141:325699
TITLE: Methods for inhibiting angiogenesis, tumor growth and metastasis by using Stq-peptides as antagonists to bind to denatured laminin
INVENTOR(S): Brooks, Peter C.; Akalu, Abebe
PATENT ASSIGNEE(S): New York University, USA
SOURCE: PCT Int. Appl., 50 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004087734	A2	20041014	WO 2004-US9332	20040326
WO 2004087734	A3	20050728		
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RW:	BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
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AU 2004225986	A1	20041014	AU 2004-225986	20040326
CA 2520372	A1	20041014	CA 2004-2520372	20040326
EP 1611151	A2	20060104	EP 2004-758409	20040326

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK
 JP 2006524241 T 20061026 JP 2006-509356 20040326
 PRIORITY APPLN. INFO.: US 2003-458523P P 20030328
 WO 2004-US9332 A 20040326

AB The invention describes methods for inhibiting angiogenesis, tumor growth and metastasis in a tissue of a mammal by administering an antagonist that specifically binds to a proteolyzed or denatured laminin with substantially greater affinity than to the native form of laminin. Methods utilizing such antagonists for therapeutic treatment of tumor growth, tumor metastasis or of restenosis also are described, as are methods to use such antagonists as diagnostic markers of angiogenesis in normal or diseased tissues both in vivo and ex vivo.

IT 771528-84-8P

RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

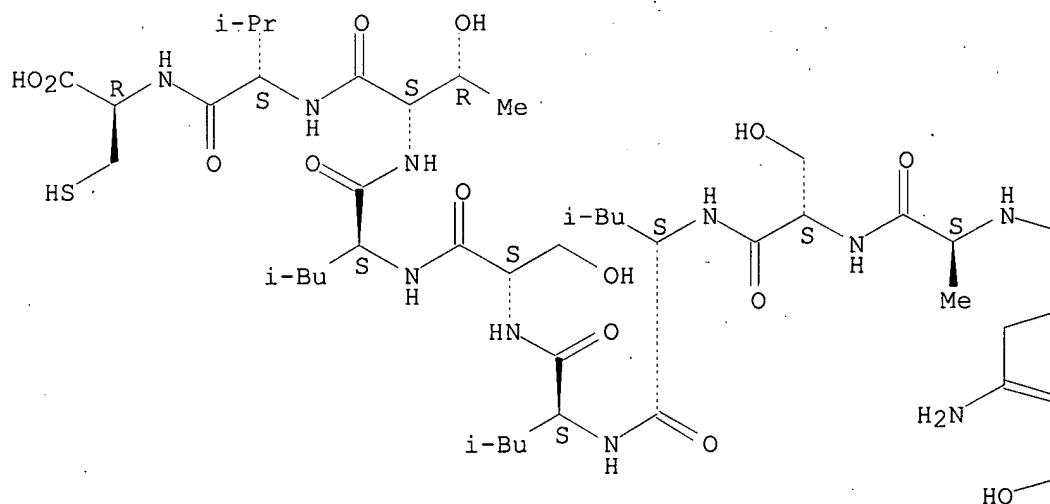
(amino acid sequence; inhibiting angiogenesis, tumor growth and metastasis by using Stq-peptides as antagonists to bind to denatured laminin)

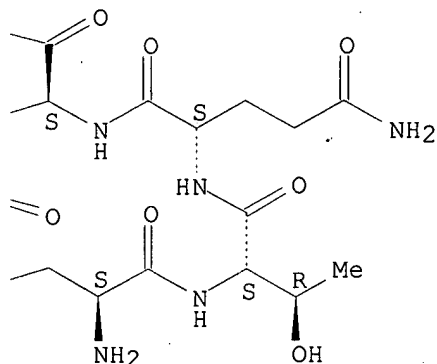
RN 771528-84-8 HCAPLUS

CN L-Cysteine, L-seryl-L-threonyl-L-glutaminyl-L-asparaginyl-L-alanyl-L-seryl-L-leucyl-L-leucyl-L-seryl-L-leucyl-L-threonyl-L-valyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A





=> FIL REG

COST IN U.S. DOLLARS

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

CA SUBSCRIBER PRICE

SINCE FILE	TOTAL
ENTRY	SESSION
15.28	44.38
SINCE FILE	TOTAL
ENTRY	SESSION
-1.50	-1.50

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T.S. Heard Ph.D.

10/797,626

=> S KGGCSTQNAQLLSLIVGKA/SQSP
L3 1 KGGCSTQNAQLLSLIVGKA/SQSP

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	29.33	73.71
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-1.50

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FILE COVERS 1907 - 16 Dec 2006 VOL 145 ISS 26
FILE LAST UPDATED: 15 Dec 2006 (20061215/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> S L3
L4 2 L3
=> D L4 1-2 IBIB ABS HITSTR

L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2006:977981 HCAPLUS
DOCUMENT NUMBER: 145:334147
TITLE: Methods of inhibiting angiogenesis and tumor development
INVENTOR(S): Brooks, Peter, C.; Akalu, Abebe; Cretu, Alexandra; Policarpio, Desiree
PATENT ASSIGNEE(S): New York University, USA
SOURCE: PCT Int. Appl., 153pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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T.S. Heard Ph.D.

WO 2006098987 A2 20060921 WO 2006-US8266 20060309

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

US 2006216236 A1 20060928 US 2006-371620 20060309

US 2006216237 A1 20060928 US 2006-372367 20060309

US 2006240002 A1 20061026 US 2006-371626 20060309

PRIORITY APPLN. INFO.: US 2005-660713P P 20050311

US 2005-660889P P 20050311

US 2005-660903P P 20050311

US 2005-711049P P 20050824

US 2005-711177P P 20050825

AB The authors disclose methods for identifying genes and proteins modulated by antagonism of extracellular matrix (ECM) ligands that specifically interact with $\alpha v \beta 3$ integrin. The authors also disclose using the identified genes and proteins for inhibiting angiogenesis, tumor metastasis, and other tumor developmental processes, including cell migration, cell adhesion, cell proliferation, and tumor growth and for treating angiogenesis-dependent conditions. In one example, a monoclonal antibody antagonist of $\alpha v \beta 3$ is shown to modulate the expression of IGFBP-4, TSP-1, Id-1, p27KIP, and p21CIP.

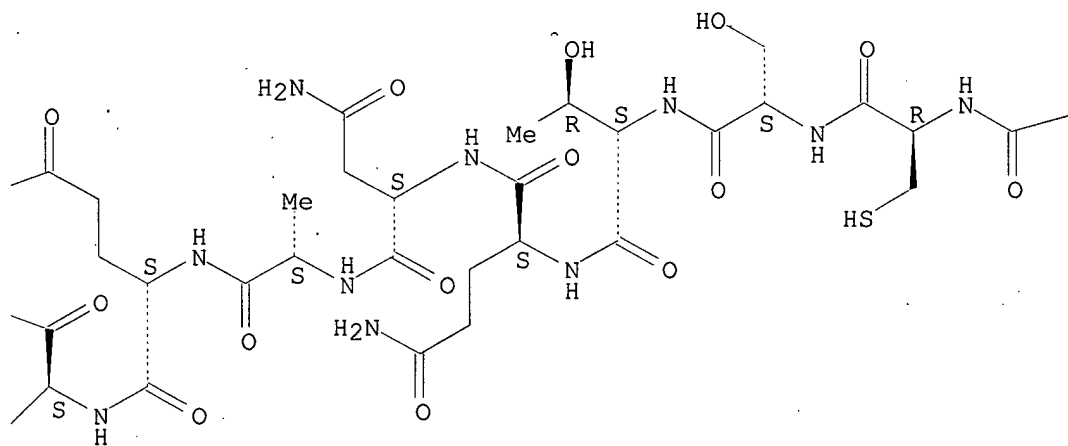
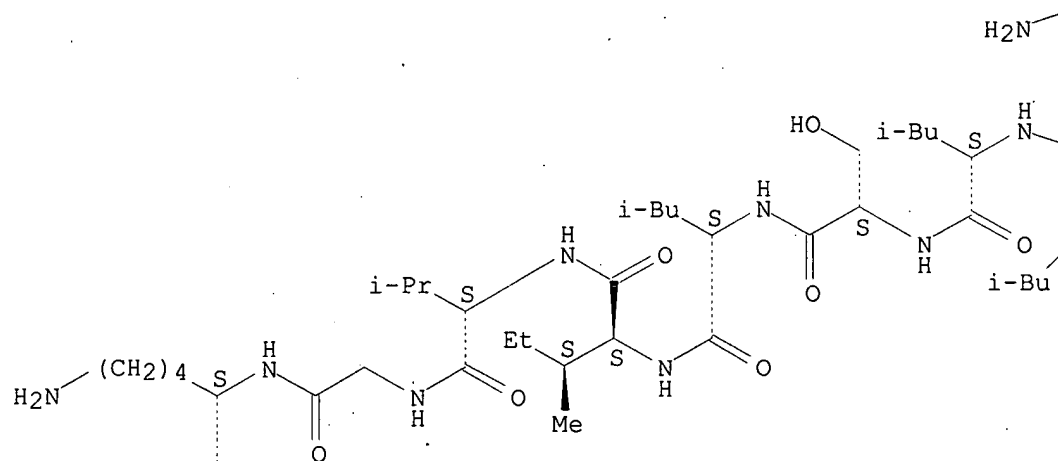
IT 771528-86-0

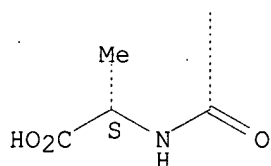
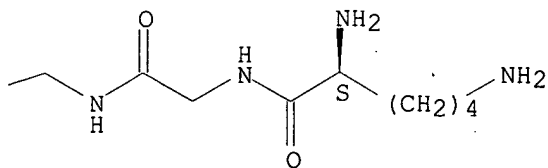
RL: PRP (Properties)
(unclaimed sequence; methods of inhibiting angiogenesis and tumor development)

RN 771528-86-0 HCAPLUS

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Absolute stereochemistry.





L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2004:857618 HCAPLUS
 DOCUMENT NUMBER: 141:325699
 TITLE: Methods for inhibiting angiogenesis, tumor growth and metastasis by using Stq-peptides as antagonists to bind to denatured laminin
 INVENTOR(S): Brooks, Peter C.; Akalu, Abebe
 PATENT ASSIGNEE(S): New York University, USA
 SOURCE: PCT Int. Appl., 50 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004087734	A2	20041014	WO 2004-US9332	20040326
WO 2004087734	A3	20050728		
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US 2004224896	A1	20041111	US 2004-797626	20040309
AU 2004225986	A1	20041014	AU 2004-225986	20040326
CA 2520372	A1	20041014	CA 2004-2520372	20040326
EP 1611151	A2	20060104	EP 2004-758409	20040326
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JP 2006524241 T 20061026 JP 2006-509356 20040326
 PRIORITY APPLN. INFO.: US 2003-458523P P 20030328
 WO 2004-US9332 A 20040326

AB The invention describes methods for inhibiting angiogenesis, tumor growth and metastasis in a tissue of a mammal by administering an antagonist that specifically binds to a proteolyzed or denatured laminin with substantially greater affinity than to the native form of laminin. Methods utilizing such antagonists for therapeutic treatment of tumor growth, tumor metastasis or of restenosis also are described, as are methods to use such antagonists as diagnostic markers of angiogenesis in normal or diseased tissues both in vivo and ex vivo.

IT 771528-86-0P

RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

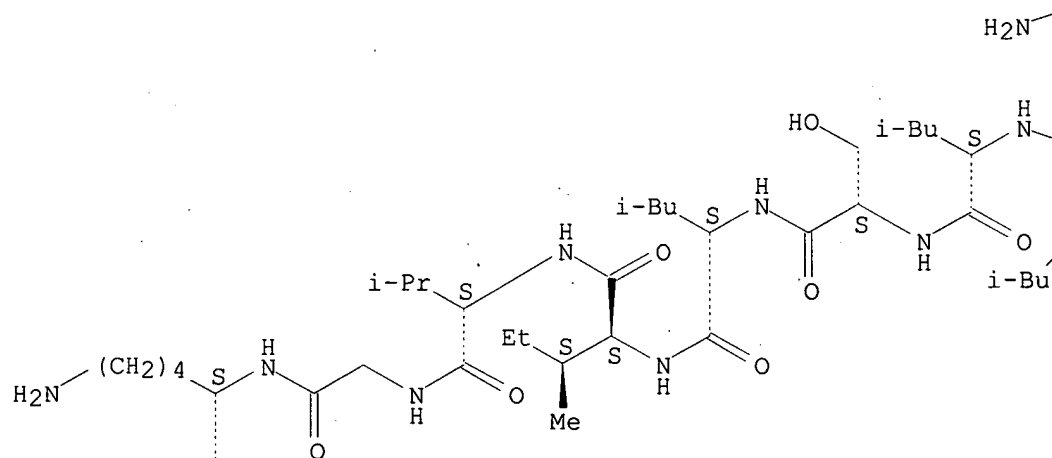
(amino acid sequence; inhibiting angiogenesis, tumor growth and metastasis by using Stq-peptides as antagonists to bind to denatured laminin)

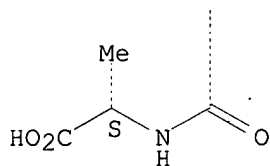
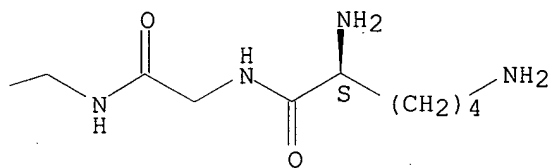
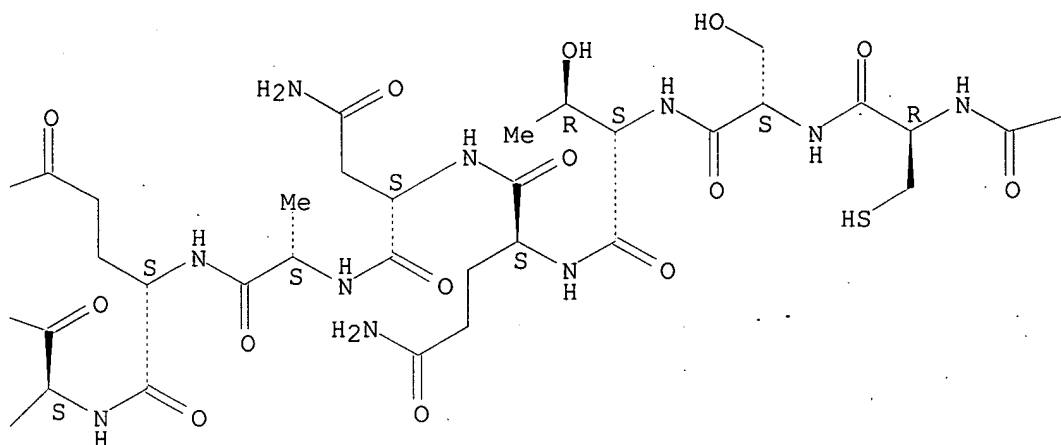
RN 771528-86-0 HCAPLUS

CN L-Alanine, L-lysylglycylglycyl-L-cysteiny-L-seryl-L-threonyl-L-glutaminyl-L-asparaginy-L-alanyl-L-glutaminyl-L-leucyl-L-leucyl-L-seryl-L-leucyl-L-isoleucyl-L-valylglycyl-L-lysyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A





=> FIL REG
COST IN U.S. DOLLARS

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

CA SUBSCRIBER PRICE

SINCE FILE	TOTAL
ENTRY	SESSION
15.28	88.99

SINCE FILE	TOTAL
ENTRY	SESSION
-1.50	-3.00

FILE 'REGISTRY' ENTERED AT 15:25:43 ON 16 DEC 2006
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STRUCTURE FILE UPDATES: 15 DEC 2006 HIGHEST RN 915749-75-6
DICTIONARY FILE UPDATES: 15 DEC 2006 HIGHEST RN 915749-75-6

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REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

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=> FIL HCAP

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
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FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

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=> S L5
L6 2 L5

T.S. Heard Ph.D.

10/797,626

=> D L6 1-2 IBIB ABS HITSTR

L6 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2006:977981 HCAPLUS

DOCUMENT NUMBER: 145:334147

TITLE: Methods of inhibiting angiogenesis and tumor development

INVENTOR(S): Brooks, Peter, C.; Akalu, Abebe; Cretu, Alexandra; Policarpio, Desiree

PATENT ASSIGNEE(S): New York University, USA

SOURCE: PCT Int. Appl., 153pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

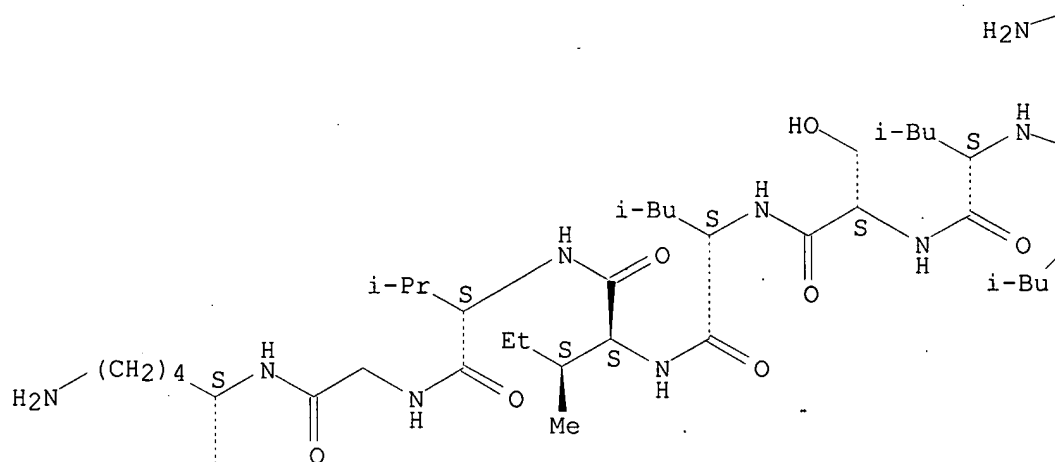
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
US 2006216236	A1	20060928	US 2006-371620	20060309
US 2006216237	A1	20060928	US 2006-372367	20060309
US 2006240002	A1	20061026	US 2006-371626	20060309
PRIORITY APPLN. INFO.:			US 2005-660713P	P 20050311
			US 2005-660889P	P 20050311
			US 2005-660903P	P 20050311
			US 2005-711049P	P 20050824
			US 2005-711177P	P 20050825
AB	The authors disclose methods for identifying genes and proteins modulated by antagonism of extracellular matrix (ECM) ligands that specifically interact with $\alpha v \beta 3$ integrin. The authors also disclose using the identified genes and proteins for inhibiting angiogenesis, tumor metastasis, and other tumor developmental processes, including cell migration, cell adhesion, cell proliferation, and tumor growth and for treating angiogenesis-dependent conditions. In one example, a monoclonal antibody antagonist of $\alpha v \beta 3$ is shown to modulate the expression of IGFBP-4, TSP-1, Id-1, p27KIP, and p21CIP.			
IT	771528-88-2			
	RL: PRP (Properties) (unclaimed sequence; methods of inhibiting angiogenesis and tumor development)			
RN	771528-88-2 HCAPLUS			
CN	L-Alanine, L-lysylglycylglycyl-L-seryl-L-threonyl-L-glutaminyl-L-asparaginyl-L-alanyl-L-glutaminyl-L-leucyl-L-leucyl-L-seryl-L-leucyl-L-isoleucyl-L-valylglycyl-L-lysyl- (9CI) (CA INDEX NAME)			

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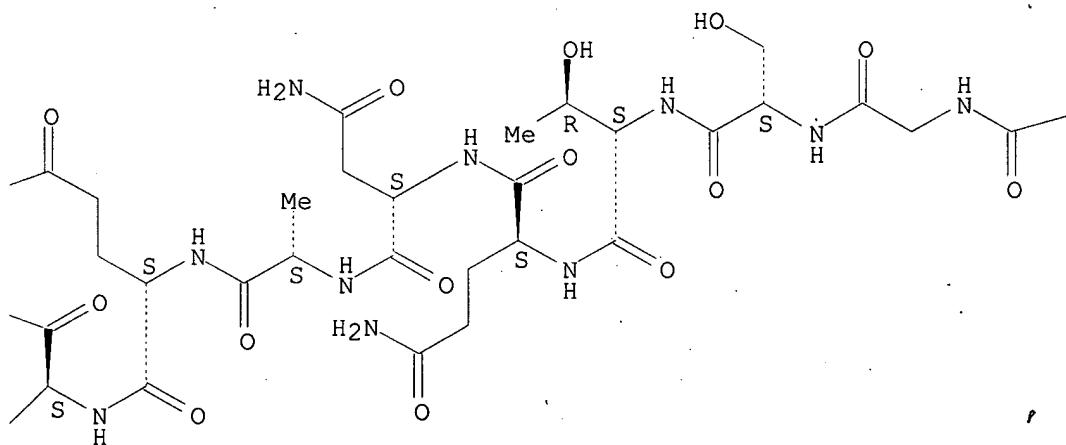
10/797,626

Absolute stereochemistry.

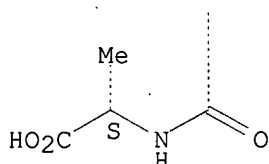
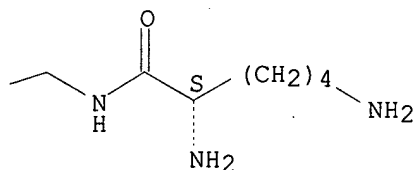
PAGE 1-A



PAGE 1-B



T.S. Heard Ph.D.



L6 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2004:857618 HCAPLUS
 DOCUMENT NUMBER: 141:325699
 TITLE: Methods for inhibiting angiogenesis, tumor growth and metastasis by using Stq-peptides as antagonists to bind to denatured laminin
 INVENTOR(S): Brooks, Peter C.; Akalu, Abebe
 PATENT ASSIGNEE(S): New York University, USA
 SOURCE: PCT Int. Appl., 50 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004087734	A2	20041014	WO 2004-US9332	20040326
WO 2004087734	A3	20050728		
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RW:	BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
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AU 2004225986	A1	20041014	AU 2004-225986	20040326
CA 2520372	A1	20041014	CA 2004-2520372	20040326
EP 1611151	A2	20060104	EP 2004-758409	20040326
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK			

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JP 2006524241	T	20061026	JP 2006-509356	20040326
PRIORITY APPLN. INFO.:			US 2003-458523P	P 20030328
			WO 2004-US9332	A 20040326

AB The invention describes methods for inhibiting angiogenesis, tumor growth and metastasis in a tissue of a mammal by administering an antagonist that specifically binds to a proteolyzed or denatured laminin with substantially greater affinity than to the native form of laminin. Methods utilizing such antagonists for therapeutic treatment of tumor growth, tumor metastasis or of restenosis also are described, as are methods to use such antagonists as diagnostic markers of angiogenesis in normal or diseased tissues both in vivo and ex vivo.

IT 771528-88-2P

RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

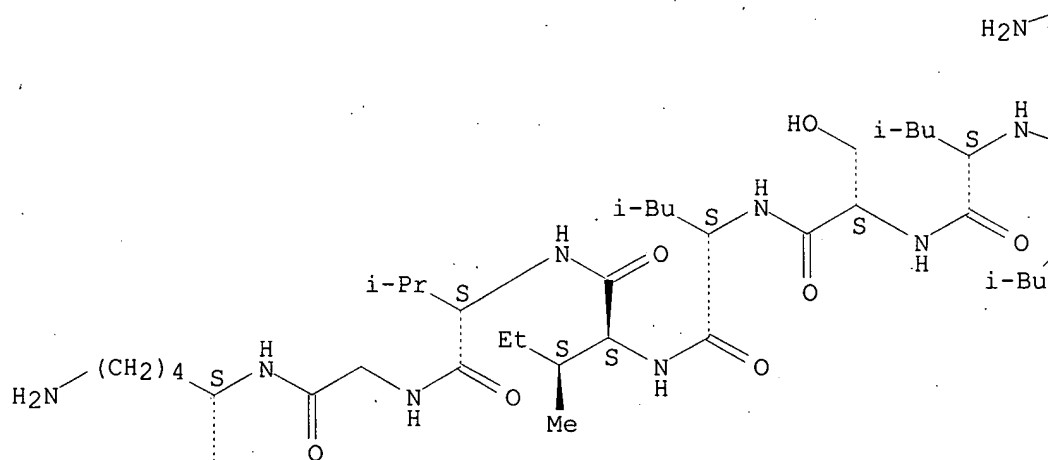
(amino acid sequence; inhibiting angiogenesis, tumor growth and metastasis by using Stq-peptides as antagonists to bind to denatured laminin)

RN 771528-88-2 HCAPLUS

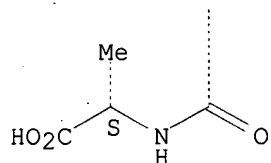
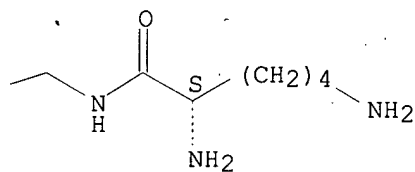
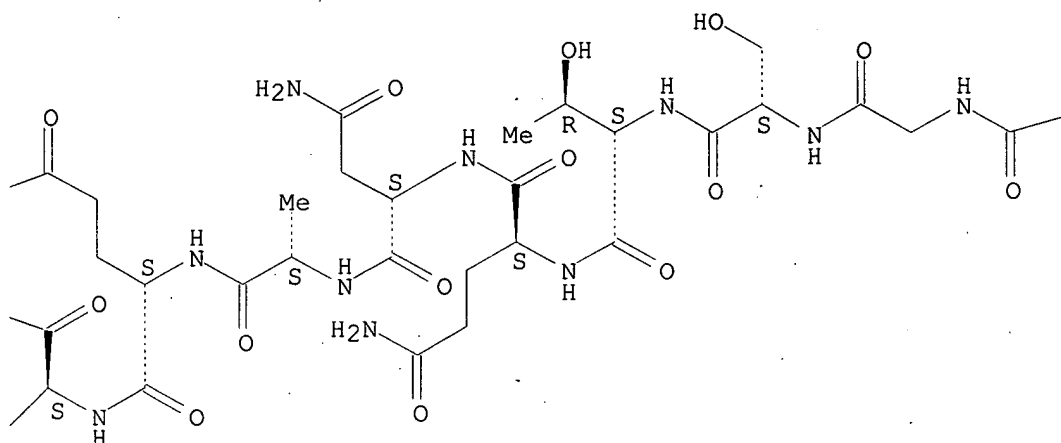
CN L-Alanine, L-lysylglycylglycyl-L-seryl-L-threonyl-L-glutaminyl-L-asparaginyl-L-alanyl-L-glutaminyl-L-leucyl-L-leucyl-L-seryl-L-leucyl-L-isoleucyl-L-valylglycyl-L-lysyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



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=> D QUE STAT

L5 1 SEA FILE=REGISTRY ABB=ON PLU=ON KGGSTQNAQLLSLIVGKA/SQSP
L6 2 SEA FILE=HCAPLUS ABB=ON PLU=ON L5

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L2 2 SEA ABB=ON PLU=ON L1
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FILE 'HCAPLUS' ENTERED AT 15:26:14 ON 16 DEC 2006

L6 2 SEA ABB=ON PLU=ON L5
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FILE HOME

FILE REGISTRY

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DICTIONARY FILE UPDATES: 15 DEC 2006 HIGHEST RN 915749-75-6

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<http://www.cas.org/ONLINE/UG/regprops.html>

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